## WHAT IS CLAIMED IS:

- 1. A substantially pure O-fucosyltransferase enzyme which is capable of glycosylating an EGF domain of a polypeptide with an activated O-fucose moiety.
- 2. The enzyme of claim 1 wherein the polypeptide glycosylated has the sequence -Cys-Xaa-Xaa-Xaa-Xaa-Ser/Thr-Cys-.
- 3. The enzyme of claim 2 wherein the polypeptide glycosylated had the sequence -Cys-Xaa-Xaa-Gly-Gly-Ser/Thr-Cys-.
- 4. The enzyme of claim 1 which has the following sequence:

  MPAGSWDPAGYLLYCPCMGRFGNQADHFLGSLAFAKLLNRTLAVPPWIEYQHHKPPFTNLH

  [SEQ ID NO:9].
- 5. The enzyme of claim 1 which has the following sequence: RLAGSWDLAGYLLYXPXMGRFGNQADHFLGSLAFAKLXVRTLAVPPWIEYQHHKPPFTNLH [SEQ ID NO:3].
- 6. A substantially pure functional fragment or analog of an O-fucosyltransferase substantially identical to the sequence:

MPAGSWDPAGYLLYCPCMGRFGNQADHFLGSLAFAKLLNRTLAVPPWIEYQHHKPPFTNLH [SEQ ID NO:9].

that is capable of glycosylating an EGF domain of a polypeptide with an activated O-fucose moiety.

- 7. The fragment or analog of claim 6 wherein the polypeptide glycosylated has the sequence -Cys-Xaa-Xaa-Xaa-Ser/Thr-Cys-.
- 8. The fragment or analog of claim 6 wherein the polypeptide glycosylated had the sequence -Cys-Xaa-Xaa-Gly-Gly-Ser/Thr-Cys-.
- 9. A substantially pure DNA sequence substantially identical to the first 1100 nucleotides of Fig. 12A [Seq. ID No. 16] which encodes a protein is capable of glycosylating the EGF domain of a polypeptide.

- 10. The DNA of claim 9 wherein the encoded protein is capable of glycosylating the sequence -Cys-Xaa-Xaa-Xaa-Xaa-Ser/Thr-Cys-.
- 11. The DNA of claim 9 wherein the encoded protein is capable of glycosylating the sequence -Cys-Xaa-Xaa-Gly-Gly-Ser/Thr-Cys-.
- 12. An O-fucosyltransferase variant which inhibits natural O-fucosyltransferase activity.
- 13. The variant of claim 12 which inhibits fucosylation of the sequence -Cys-Xaa-Xaa-Xaa-Xaa-Ser/Thr-Cys-.
- 14. The variant of claim 12 which inhibits fucosylation of the sequence -Cys-Xaa-Xaa-Gly-Gly-Ser/Thr-Cys-.
- 15. An antibody which is capable of binding O-fucosyltransferase.
- 16. The antibody of claim 15 which is capable of binding O-fucosylatransferase of the sequence: MPAGSWDPAGYLLYCPCMGRFGNQADHFLGSLAFAKLLNRTLAVPPWIEYQHHKPPFTNLH [SEQ ID NO:9].
- 17. The antibody of claim 15 which is polyclonal.
- 18. The antibody of claim 15 which is monoclonal.
- 19. The antibody of claim 15 which is humanized.
- 20. The antibody of claim 15 which is bispecific.
- 21. The antibody of claim 15 which is heterospecific.
- 22. A method of glycosylating a EGF domain of a polypeptide with an activated O-fucose moiety comprising the application of an O-fucosyltransferase.
- 23. The method of claim 22 wherein the glycosylated polypeptide comprises the sequence -Cys-Xaa-Xaa-Xaa-Xaa-Ser/Thr-Cys-.

24. The method of claim 22 wherein the glycosylated polypeptide comprises the sequence -Cys-Xaa-Xaa-Gly-Gly-Ser/Thr-Cys-.

- 25. A process for isolating and purifying O-fucosyltransferase comprising:
  - a) preparing an extract from a cell line expressing O-fucosyltransferase,
  - b) purifying via a first chromatography step over sequentially applied anion exchange resin and nucleotide binding resin;
  - c) purifying via a second chromatography step over an acceptor substrate ligand associated with a metal chelating-agarose resin;
  - d) purifying via a third chromatography purification over a donor substrate analog ligand associated with agarose.
- 26. The process of claim 25 wherein the anion exchange resin is DE-52.
- 27. The process of claim 25 wherein the nucleotide binding resin is Cibacron Blue 3GA.
- 28. The process of claim 25 wherein the donor substrate analog is GDP-hexanolamine.
- 29. The process of claim 25 wherein the metal chelating resin is an IMAC resin.
- 30. The process of claim 25 wherein the metal chelating resin is Ni<sup>2+</sup>-NTA.